

Hydrocortisel

Authorised

- NEOMYCIN SULFATE
- Dexamethasone acetate

Product identification

Medicine name:

Hydrocortisel

Active substance:

NEOMYCIN SULFATE

Dexamethasone acetate

Target species:

Dog

Route of administration:

Cutaneous use

Product details

Active substance and strength:

NEOMYCIN SULFATE

12.00 milligram(s) / 1.00 millilitre(s)

Dexamethasone acetate

0.17 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Cutaneous emulsion

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD07CB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Package description:

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Selectavet Dr. Otto Fischer GmbH

Marketing authorisation date:

30/11/2004

Manufacturing sites for batch release:

C.P.M. ContractPharma GmbH

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

400688.00.00

Date of authorisation status change:

17/09/2012

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.